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10/734,272	12/15/2003	Yu-Fang Hu	MR929-944	1490

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EXAMINER

KISHORE, GOLLAMUDI S

ART UNIT	PAPER NUMBER
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1612

NOTIFICATION DATE	DELIVERY MODE
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07/16/2010

ELECTRONIC

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

ptoactions@rklpatlaw.com
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DETAILED ACTION

The amendment dated 6-6-08 is acknowledged.

Claims included in the prosecution are 1-24 and 26-30.

In view of the amendments the 102 rejection is withdrawn. In view of the cancellation of claim 25, the 102 rejection over Slater, Zalipsky and Papahadjopoulos is withdrawn.

Claim Rejections - 35 USC § 112

1. The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

2. Claims 1-24 and 26-30 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Applicant amends claim 1 to introduce the expression, "for increasing extrusion speed and lowering extrusion pressure". It is unclear as to how much the speed is increased to and the pressure is lowered to.

Claim Rejections - 35 USC § 103

3. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Art Unit: 1612

4. Claims 1-24 and 26-30 are rejected under 35 U.S.C. 103(a) as being unpatentable over Slater (6,355,268) by itself or in combination with Zalipsky (6,051,251).

Slater discloses a process of preparation of liposomes. The process involves mixing soy phosphatidylcholine, cholesterol and PEG-DSPE in a mole ratio of 56.4:38.3:5.3 in ethanol at 65 degrees and mixing this mixture with an ammonium sulfate solution at 65 degrees. The mixture is subjected to an extrusion process through an extruder. The ammonium sulfate and ethanol are removed from the external bulk aqueous phase prior to loading the active agent. The active agent, Topotecan is dissolved in 40 ml of 10 % sucrose solution and then remotely loaded. The method of Slater differs from instant method in that, the ammonium sulfate and ethanol are removed by diafiltration and not by dialysis process (abstract, col. 7, line 20 through col. 9, line 23 and Example 1). One could however, interpret diafiltration as a dialysis process. Since the goal is to remove ammonium sulfate and ethanol from the liposomal mixture, it would have been obvious to one of ordinary skill in the art to use art known process of dialysis, which removes salts with a reasonable expectation of success. One of ordinary skill in the art would be motivated to use this process since the reference of Zalipsky shows the removal of ammonium sulfate from the external medium by dialysis and also equates dialysis and diafiltration for the removal of small molecular weight compounds such as drugs (Example 1 and col. 8, lines 12-16). Although Slater does not disclose claimed ranges for the individual components, since both instant invention and Slater are involved in the process of preparation of liposomes loaded with the active

Art Unit: 1612

agents, it is deemed obvious to one of ordinary skill in the art to manipulate the amounts of the phospholipids and ethanol to obtain the best possible results. Slater does not disclose the active agent to be doxorubicin. However, since the principle of loading the active agent is the same, one of ordinary skill in the art would be able to load any active agent in the method of Slater with a reasonable expectation of success. Although Slater does not teach lyophilization of liposomes, since this step is routinely practiced in the art of liposomes, one of ordinary skill in the art would be motivated to lyophilize the liposomes with a reasonable expectation of success.

Applicant's arguments have been fully considered, but are not persuasive.

Applicant argues that the main objective of the Slater reference is to provide a topoisomerase inhibitor composition to improve cancer therapy and to provide a liposome composition for administration of the topoisomerase inhibitor and not about a preparing method of liposome and merely comprises a conventional method of preparation. This argument is not persuasive since irrespective of the main objective, Slater discloses a method which is similar to instant method. Applicant argues that extrusion process of the liposome preparation usually experiences difficulties at high extrusion pressure and low extrusion speed and that the present invention provides a production process of liposome suspension with low pressure and a higher extrusion speed. Applicant further argues that the present invention discloses a proper amount of mixture, the alcohol solvent and the ammonium sulfate solution and that the ratio of the alcohol and the mixture does not fall within the ratio range of the present invention.

These arguments are not persuasive since it is within the skill of the art to recognize

Art Unit: 1612

that more the viscous to solution is, the more pressure and speed are required for extrusion. The examiner cites in this context, the reference of Schneider (5,393,530) which teaches the connection between the viscosity and extrusion process. On col. 7, lines 19-21 Schneider states "Also extrusion of empty liposomes is easy because of their inherent low viscosity". This statement implies a connection between the viscosity of the solution and the extrusion process. The examiner also cites the references of Suddith which teaches fluid viscosity affects the extrusion rate (col. 6, lines 8-14). Therefore, it would have been obvious to one of ordinary skill in the art to increase the amounts of alcohol if necessary, to reduce the viscosity of the lipid solution in ethanol. Instant invention therefore, is an obvious extension of Slater's method.

Applicant argues that Zalipsky reference does not overcome the deficiencies of Slater since the objective of Zalipsky is not about the liposome preparation process. This argument is not persuasive since irrespective of the objective, Zalipsky is combined for its teachings of the removal of ammonium sulfate from the external medium by dialysis and for its teachings of equivalency between dialysis and diafiltration for the removal of small molecular weight compounds such as drugs.

5. Claims 1-24 and 26-30 are rejected under 35 U.S.C. 103(a) as being unpatentable over Slater (6,355,268) by itself or in combination with Zalipsky (6,051,251) as set forth above, further in view of Leigh (5,004,611) and Suddith (5,556,580).

The teachings of Slater and Zalipsky have been discussed above.

Art Unit: 1612

Leigh while disclosing proliposomal formulations teaches that the ratio of the lipid component to water-miscible component (ethanol) could be from 1:2 to 1:10 (abstract, col. 4, lines 35-38).

Suddith while disclosing an extrusion process of liposomes teaches that the viscosity affects the extrusion rate (col. 2, lines 52-55 and col. 6, lines 8-14).

It would have been obvious to one of ordinary skill in the art to vary the amounts of alcohol in lipid solutions of Slater to reduce their viscosity in order to facilitate extrusion at higher speed and lower pressure, if such are desired, since Leigh teaches that alcohol can be increased to 10:1 compared to the lipid amount and that of Suddith teaches that viscosity of the solution affects the extrusion rate.

6. Claims 27-29 are rejected under 35 U.S.C. 103(a) as being unpatentable over Slater (6,355,268) by itself or in combination with Zalipsky (6,051,251) OR over Slater (6,355,268) by itself or in combination with Zalipsky (6,051,251) as set forth above, further in view of Leigh (5,004,611) and Suddith (5,556,580) both set forth above, further in view of Barenholz (5,316,771).

The teachings of Slater, Zalipsky, Leigh and Suddith have been discussed above. What is lacking in these references is the teaching of doxorubicin as the active agent. The use of doxorubicin as the active agent in the liposomes of Slater would have been obvious to one of ordinary skill in the art with a reasonable expectation of success since the reference of Barenholz shows the loading of doxorubicin in liposomes containing ammonium sulfate (fig. 3, examples and claims).

Applicant's arguments have been fully considered, but are not persuasive.

Applicant argues that Barenholz focuses on the drug loading method in liposomes and that Barenholz never addresses the high pressure and the low extrusion speed caused during the extrusion process. This argument is not persuasive since this reference is combined for its teachings of the encapsulation of doxorubicin in liposomes and not for its teachings or lack thereof of extrusion process.

7. Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to GOLLAMUDI S. KISHORE whose telephone number is (571)272-0598. The examiner can normally be reached on 6:30 AM- 4 PM, alternate Friday off.

Art Unit: 1612

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Krass Frederick can be reached on (571) 272-0580. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Gollamudi S Kishore/
Primary Examiner, Art Unit 1612

GSK